Peter R Shepherd

List of Publications by Year
in descending order

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78 papers	5,264 citations	293460 24 h-index	93651 72 g-index
80 all docs	80 docs citations	80 times ranked	9445 citing authors

#	Article	IF	Citations
1	Derivation of induced pluripotent stem cell lines from New Zealand donors. Journal of the Royal Society of New Zealand, 2022, 52, 54-67.	1.0	5
2	The Impact of Exogenous Insulin Input on Calculating Hepatic Clearance Parameters. Journal of Diabetes Science and Technology, 2022, 16, 945-954.	1.3	4
3	The minor allele of the CREBRF rs373863828 p.R457Q coding variant is associated with reduced levels of myostatin in males: Implications for body composition. Molecular Metabolism, 2022, 59, 101464.	3.0	2
4	Î²-Cells retain a pool of insulin-containing secretory vesicles regulated by adherens junctions and the cadherin-binding protein p120 catenin. Journal of Biological Chemistry, 2022, 298, 102240.	1.6	0
5	Genomic and signalling pathway characterization of the NZM panel of melanoma cell lines: A valuable model for studying the impact of genetic diversity in melanoma. Pigment Cell and Melanoma Research, 2021, 34, 136-143.	1.5	9
6	Efficacy of Providing the PI3K p110Î± Inhibitor BYL719 (Alpelisib) to Middle-Aged Mice in Their Diet. Biomolecules, 2021, 11, 150.	1.8	6
7	Diverse mechanisms activate the PI 3-kinase/mTOR pathway in melanomas: implications for the use of PI 3-kinase inhibitors to overcome resistance to inhibitors of BRAF and MEK. BMC Cancer, 2021, 21, 136.	1.1	21
8	Variability in Estimated Modelled Insulin Secretion. Journal of Diabetes Science and Technology, 2021, , 193229682199112.	1.3	2
9	A role for PAK1 mediated phosphorylation of Î²-catenin Ser552 in the regulation of insulin secretion. Biochemical Journal, 2021, 478, 1605-1615.	1.7	6
10	Glucose regulates expression of pro-inflammatory genes, IL-1Î² and IL-12, through a mechanism involving hexosamine biosynthesis pathway-dependent regulation of Î±-E catenin. Bioscience Reports, 2021, 41, .	1.1	7
11	Î²â€Catenin is required for optimal exerciseâ€•and contractionâ€stimulated skeletal muscle glucose uptake. Journal of Physiology, 2021, 599, 3897-3912.	1.3	12
12	The CREBRF diabetes-protective rs373863828-A allele is associated with enhanced early insulin release in men of MÄori and Pacific ancestry. Diabetologia, 2021, 64, 2779-2789.	2.9	7
13	The CSF1 receptor inhibitor pexidartinib (PLX3397) reduces tissue macrophage levels without affecting glucose homeostasis in mice. International Journal of Obesity, 2020, 44, 245-253.	1.6	39
14	The MÄori and Pacific specific CREBRF variant and adult height. International Journal of Obesity, 2020, 44, 748-752.	1.6	15
15	Î²-catenin regulates muscle glucose transport via actin remodelling and M-cadherin binding. Molecular Metabolism, 2020, 42, 101091.	3.0	26
16	Cetuximab produced from a goat mammary gland expression system is equally efficacious as innovator cetuximab in animal cancer models. Biotechnology Reports (Amsterdam, Netherlands), 2020, 28, e00533.	2.1	2
17	The Potential of Anti-Diabetic RÄkau RongoÄ•(MÄori Herbal Medicine) to Treat Type 2 Diabetes Mellitus (T2DM) Mate Huka: A Review. Frontiers in Pharmacology, 2020, 11, 935.	1.6	6
18	Î±-catenin isoforms are regulated by glucose and involved in regulating insulin secretion in rat clonal Î²-cell models. Biochemical Journal, 2020, 477, 763-772.	1.7	8

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19	Prolonged treatment with a PI3K p110Î± inhibitor causes sex- and tissue-dependent changes in antioxidant content, but does not affect mitochondrial function. Bioscience Reports, 2020, 40, .	1.1	6
20	Evolution of Molecular Targets in Melanoma Treatment. Current Pharmaceutical Design, 2020, 26, 396-414.	0.9	10
21	For Better or Worse: The Potential for Dose Limiting the On-Target Toxicity of PI 3-Kinase Inhibitors. Biomolecules, 2019, 9, 402.	1.8	16
22	Synthesis and Evaluation of Imidazo[1,2â€‹a]pyridine Analogues of the ZSTK474 Class of Phosphatidylinositol 3â€Kinase Inhibitors. Chemistry - an Asian Journal, 2019, 14, 1249-1261.	1.7	9
23	Orthogonal assays for the identification of inhibitors of the single-stranded nucleic acid binding protein YB-1. Acta Pharmaceutica Sinica B, 2019, 9, 997-1007.	5.7	6
24	Synthesis and biological evaluation of solubilized sulfonamide analogues of the phosphatidylinositol 3-kinase inhibitor ZSTK474. Bioorganic and Medicinal Chemistry, 2019, 27, 1529-1545.	1.4	12
25	The role of adherens junction proteins in the regulation of insulin secretion. Bioscience Reports, 2018, 38, .	1.1	29
26	Î²-catenin is important for the development of an insulin responsive pool of GLUT4 glucose transporters in 3T3-L1 adipocytes. Experimental Cell Research, 2018, 366, 49-54.	1.2	17
27	Macrophages enhance Vegfa-driven angiogenesis in an embryonic zebrafish tumour xenograft model. DMM Disease Models and Mechanisms, 2018, 11, .	1.2	47
28	Re: â€œWidespread prevalence of a CREBRF variant among MÄori and Pacific children is associated with weight and height in early childhoodâ€• International Journal of Obesity, 2018, 42, 1389-1391.	1.6	5
29	Feeding and <scp>glucagonâ€‘like peptide</scp>â€‘1 receptor activation stabilise Î²â€‘catenin in specific hypothalamic nuclei in male rats. Journal of Neuroendocrinology, 2018, 30, e12607.	1.2	4
30	Discordant association of the CREBRF rs373863828 A allele with increased BMI and protection from type 2 diabetes in MÄori and Pacific (Polynesian) people living in Aotearoa/New Zealand. Diabetologia, 2018, 61, 1603-1613.	2.9	61
31	Niclosamide reduces glucagon sensitivity via hepatic PKA inhibition in obese mice: Implications for glucose metabolism improvements in type 2 diabetes. Scientific Reports, 2017, 7, 40159.	1.6	23
32	Novel pyrazolo[1,5- a]pyridines with improved aqueous solubility as p110Î±-selective PI3 kinase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2017, 27, 187-190.	1.0	9
33	Combining properties of different classes of PI3KÎ± inhibitors to understand the molecular features that confer selectivity. Biochemical Journal, 2017, 474, 2261-2276.	1.7	6
34	Synthesis and biological evaluation of sulfonamide analogues of the phosphatidylinositol 3-kinase inhibitor ZSTK474. Bioorganic and Medicinal Chemistry, 2017, 25, 5859-5874.	1.4	14
35	Biological characterization of SN32976, a selective inhibitor of PI3K and mTOR with preferential activity to PI3KÎ±, in comparison to established pan PI3K inhibitors. Oncotarget, 2017, 8, 47725-47740.	0.8	11
36	Niclosamide blocks glucagon phosphorylation of Ser552 on Î²-catenin in primary rat hepatocytes via PKA signalling. Biochemical Journal, 2016, 473, 1247-1255.	1.7	19

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37	A Critical Role for Î²-Catenin in Modulating Levels of Insulin Secretion from Î²-Cells by Regulating Actin Cytoskeleton and Insulin Vesicle Localization. Journal of Biological Chemistry, 2016, 291, 25888-25900.	1.6	42
38	Targeted inhibition of dominant PI3-kinase catalytic isoforms increase expression of stem cell genes in glioblastoma cancer stem cell models. International Journal of Oncology, 2016, 49, 207-216.	1.4	13
39	Inhibitors of pan-PI3K Signaling Synergize with BRAF or MEK Inhibitors to Prevent BRAF-Mutant Melanoma Cell Growth. Frontiers in Oncology, 2015, 5, 135.	1.3	52
40	Glucagon phosphorylates serine 552 ofÎ²-catenin leading to increased expression of cyclin D1 and c-Myc in the isolated rat liver. Archives of Physiology and Biochemistry, 2015, 121, 88-96.	1.0	13
41	Exploring the isoform selectivity of TGX-221 related pyrido[1,2-a]pyrimidinone-based Class IA PI 3-kinase inhibitors: Synthesis, biological evaluation and molecular modelling. Bioorganic and Medicinal Chemistry, 2015, 23, 3796-3808.	1.4	9
42	Novel pyrazolo[1,5-a]pyridines as PI3K inhibitors: variation of the central linker group. MedChemComm, 2014, 5, 41-46.	3.5	12
43	Structure, function and inhibition of the phosphoinositide 3-kinase p110Î± enzyme. Biochemical Society Transactions, 2014, 42, 120-124.	1.6	11
44	Targeting Class IA PI3K Isoforms Selectively Impairs Cell Growth, Survival, and Migration in Glioblastoma. PLoS ONE, 2014, 9, e94132.	1.1	33
45	Extended treatment with selective phosphatidylinositol 3â€kinase and m<scp>TOR</scp> inhibitors has effects on metabolism, growth, behaviour and bone strength. FEBS Journal, 2013, 280, 5337-5349.	2.2	22
46	Enzyme activity effects of N-terminal His-tag attached to catalytic sub-unit of phosphoinositide-3-kinase. Bioscience Reports, 2013, 33, .	1.1	20
47	Identification of a pathway by which glucose regulates Î²-catenin signalling via the cAMP/protein kinase A pathway in Î²-cell models. Biochemical Journal, 2013, 449, 803-811.	1.7	25
48	Oncogenic Mutations of p110Î± Isoform of PI 3-Kinase Upregulate Its Protein Kinase Activity. PLoS ONE, 2013, 8, e71337.	1.1	14
49	Î°B kinase Î² (IKKÎ²) does not mediate feedback inhibition of the insulin signalling cascade. Biochemical Journal, 2012, 442, 723-732.	1.7	5
50	Effects of acutely inhibiting PI3K isoforms and mTOR on regulation of glucose metabolism in vivo. Biochemical Journal, 2012, 442, 161-169.	1.7	42
51	Beta-Testing of PI3-Kinase Inhibitors: Is Beta Better?. Cancer Discovery, 2012, 2, 393-394.	7.7	3
52	A drug targeting only p110Î± can block phosphoinositide 3-kinase signalling and tumour growth in certain cell types. Biochemical Journal, 2011, 438, 53-62.	1.7	137
53	Phosphoinositide 3-Kinase (PI3K(p110Î±)) Directly Regulates Key Components of the Z-disc and Cardiac Structure*. Journal of Biological Chemistry, 2011, 286, 30837-30846.	1.6	32
54	NVP-BEZ235, a dual pan class I PI3 kinase and mTOR inhibitor, promotes osteogenic differentiation in human mesenchymal stromal cells. Journal of Bone and Mineral Research, 2010, 25, 2126-2137.	3.1	54

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55	Evidence for a role for the p110-Î± isoform of PI3K in skeletal function. Biochemical and Biophysical Research Communications, 2010, 391, 564-569.	1.0	11
56	Anti-Leukemic Activity of PIK-75, a PI3-Kinase p110Î± Selective Inhibitor, In Acute Myeloid Leukemia. Blood, 2010, 116, 659-659.	0.6	0
57	Investigating the role of class-IA PI 3-kinase isoforms in adipocyte differentiation. Biochemical and Biophysical Research Communications, 2009, 379, 830-834.	1.0	15
58	Functional differences between two classes of oncogenic mutation in the PIK3CA gene. Biochemical and Biophysical Research Communications, 2009, 381, 577-581.	1.0	50
59	The Synthesis of Phosphopeptides Using Microwave-assisted Solid Phase Peptide Synthesis. International Journal of Peptide Research and Therapeutics, 2008, 14, 387-392.	0.9	29
60	Glucose induces an autocrine activation of the Wnt/Î²-catenin pathway in macrophage cell lines. Biochemical Journal, 2008, 416, 211-218.	1.7	74
61	The Role of Phosphoinositide 3-Kinase C2Î± in Insulin Signaling. Journal of Biological Chemistry, 2007, 282, 28226-28236.	1.6	136
62	Evidence for functional redundancy of class IA PI3K isoforms in insulin signalling. Biochemical Journal, 2007, 404, 449-458.	1.7	188
63	PI 3-kinase p110Î²: a new target for antithrombotic therapy. Nature Medicine, 2005, 11, 507-514.	15.2	555
64	Letter fromâ€¦ New Zealand: Doing biochemistry down under. Biochemist, 2005, 27, 43-44.	0.2	0
65	Regulation of Phosphoinositide 3-Kinase by Its Intrinsic Serine Kinase Activity In Vivo. Molecular and Cellular Biology, 2004, 24, 966-975.	1.1	60
66	eIF4E binding protein 1 and H-Ras are novel substrates for the protein kinase activity of class-I phosphoinositide 3-kinase. Biochemical and Biophysical Research Communications, 2004, 319, 541-549.	1.0	16
67	Secrets of insulin and IGF-1 regulation of insulin secretion revealed. Biochemical Journal, 2004, 377, e1-e2.	1.7	7
68	Comparison of the kinetic properties of the lipid- and protein-kinase activities of the p110Î± and p110Î² catalytic subunits of class-Ia phosphoinositide 3-kinases. Biochemical Journal, 2000, 350, 353.	1.7	20
69	Comparison of the kinetic properties of the lipid- and protein-kinase activities of the p110Î± and p110Î² catalytic subunits of class-Ia phosphoinositide 3-kinases. Biochemical Journal, 2000, 350, 353-359.	1.7	60
70	Glucose Transporters and Insulin Action â€" Implications for Insulin Resistance and Diabetes Mellitus. New England Journal of Medicine, 1999, 341, 248-257.	13.9	1,123
71	Mammalian target of rapamycin is a direct target for protein kinase B: identification of a convergence point for opposing effects of insulin and amino-acid deficiency on protein translation. Biochemical Journal, 1999, 344, 427-431.	1.7	795
72	Insulin-like Growth Factors Require Phosphatidylinositol 3-Kinase to Signal Myogenesis: Dominant Negative p85 Expression Blocks Differentiation of L6E9 Muscle Cells. Molecular Endocrinology, 1998, 12, 66-77.	3.7	98

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73	Phosphoinositide 3-kinase: the key switch mechanism in insulin signalling. Biochemical Journal, 1998, 333, 471-490.	1.7	924
74	Over-expression of GLUT4 selectively in adipose tissue in transgenic mice: Implications for nutrient partitioning. Proceedings of the Nutrition Society, 1996, 55, 191-199.	0.4	29
75	Phorbol esters stimulate phosphatidylinositol 3,4,5-trisphosphate production in 3T3-L1 adipocytes: implications for stimulation of glucose transport. Biochemical Journal, 1996, 318, 203-205.	1.7	33
76	Effect of phorbol esters on phosphatidylinositol 3-kinase activity in adipocytes. Biochemical Society Transactions, 1995, 23, 183S-183S.	1.6	4
77	Involvement of PI 3-kinase in stimulation of glucose transport and recruitment of transferrin receptors in 3T3-L1 adipocytes. Biochemical Society Transactions, 1995, 23, 201S-201S.	1.6	8
78	Insulin activates Glycogen Synthase by a novel PI 3-kinase/p70s6k dependent pathway in 3T3-L1 adipocytes. Biochemical Society Transactions, 1995, 23, 202S-202S.	1.6	8